

AH-130 FN Cells | 500451**General information****Description**

AH-130 FN is a variant of the AH-130 rat ascites tumor cell line, used extensively in studies related to coagulation, fibrinolysis, and metastasis. These cells were derived from rats and are typically maintained by serial intraperitoneal implantation in male Donryu rats. The AH-130 line itself is known for its high thromboplastic and fibrinolytic activities, which are linked to its role in promoting blood-borne metastasis, especially in the lungs. In contrast, the AH-130 FN variant has lower thromboplastic and fibrinolytic activity. This difference in enzymatic activity between AH-130 and AH-130 FN is crucial as it influences the formation of thrombi and the number of metastatic foci in the lungs following intravenous inoculation.

Research has shown that after intravenous injection, AH-130 cells cause a significant reduction in platelet count and fibrinogen levels, indicative of increased thrombus formation. This effect is notably more pronounced than in AH-130 FN. Histological studies demonstrate that AH-130 forms more abundant metastatic foci in the lungs compared to AH-130 FN, both at 72 hours and 7 days post-inoculation. AH-130 is associated with the formation of thrombi composed of platelets and fibrin around embolized tumor cells, whereas AH-130 FN shows sparse thrombus formation. These findings suggest that AH-130's higher thromboplastic activity plays a significant role in promoting metastasis through platelet aggregation and fibrin deposition around the tumor cells, a process less prominent in AH-130 FN.

Organism

Rat

Tissue

Liver

Disease

Hepatocellular carcinoma

Synonyms

AH130FN-TC, AH130FN, AH-130F(N), AH-130FN, AH 130 FN

Characteristics**Morphology**

Round cells in suspension, Epithelial-like when adherent

Growth properties

Suspension, few adherent

Regulatory Data**Citation**

AH-130 FN (Cytion catalog number 500451)

Biosafety level

1

NCBI_TaxID

10116

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CellosaurusAccession CVCL_5683

Biomolecular Data

Tumorigenic Yes, in Wistar rats.**Viruses** RAP-test negative. .

Handling

Culture Medium DMEM:Ham's F12 (1:1), w: 3.1 g/L Glucose, w: 2.5 mM L-Glutamine, w: 15 mM HEPES, w: 0.5 mM Sodium pyruvate, w: 1.2 g/L NaHCO₃ (Cytion article number 820400a)**Supplements** Supplement the medium with 10% FBS**Subculturing** Gently homogenize the cell suspension in the flask by pipetting up and down, then take a representative sample to determine the cell density per ml. Dilute the suspension to achieve a cell concentration of 1×10^5 cells/ml with fresh culture medium, and aliquot the adjusted suspension into new flasks for further cultivation.**Seeding density** 1×10^6 cells/cm²**Fluid renewal** Every 3 to 5 days**Post-Thaw Recovery** After thawing, plate the cells at 5×10^4 cells/cm² and allow the cells to recover from the freezing process and to adhere for at least 24 hours.**Freeze medium** As a cryopreservation medium, we use complete growth medium (including FBS) + 10% DMSO for adequate post-thaw viability, or CM-1 (Cytion catalog number 800100), which includes optimized osmoprotectants and metabolic stabilizers to enhance recovery and reduce cryo-induced stress.

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Thawing and Culturing Cells

1. Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
2. Upon receipt, either store the cryovial immediately at temperatures below -150°C to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required.
3. For immediate culturing, swiftly thaw the vial by immersing it in a 37°C water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains.
4. Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
5. Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently.
6. Centrifuge the mixture at 300 x g for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium.
7. Gently resuspend the cell pellet in 10 ml of fresh culture medium. For adherent cells, divide the suspension between two T25 culture flasks; for suspension cultures, transfer all the medium into one T25 flask to promote effective cell interaction and growth.
8. Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.

Incubation Atmosphere

37°C, 5% CO₂, humidified atmosphere.

Shipping Conditions

Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately -78 °C throughout transit. On receipt, inspect the container immediately and transfer vials without delay to appropriate storage.

Storage Conditions

For long-term preservation, place vials in vapor-phase liquid nitrogen at about -150 to -196 °C. Storage at -80 °C is acceptable only as a short interim step before transfer to liquid nitrogen.

Quality Control & Molecular Analysis

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Sterility

Mycoplasma contamination is excluded using both PCR-based assays and luminescence-based mycoplasma detection methods.

To ensure there is no bacterial, fungal, or yeast contamination, cell cultures are subjected to daily visual inspections.